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# In the Claims

- 1. Cancel.
- 2. Cancel.
- 3. Cancel.
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- 5. Cancel.
- 6. Cancel.
- 7.. Cancel.
- 8. Cancel.
- 9. Cancel.
- 10. Cancel.
- 11. Cancel.
- 12. Cancel.
- 13. Cancel.
- 14. Cancel.
- 15. Cancel.
- 16. Cancel.

17(Original) A compound of structural formula I:

1

or a pharmaceutically acceptable salt, enantiomer, diastereomer, pro drug or mixture thereof, wherein

X is (CH2)n, O or S;

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Ar<sub>2</sub> independently represent (CH<sub>2</sub>)<sub>m</sub>C<sub>6-10</sub>aryl, (CH<sub>2</sub>)<sub>m</sub>C<sub>5-10</sub>heteroaryl, (CH<sub>2</sub>)<sub>m</sub>C<sub>3-10</sub> heterocycloalkyl, (CH<sub>2</sub>)<sub>m</sub>C<sub>3-8</sub> cycloalkyl said cycloalkyl, heterocycloalkyl, aryl or heteroaryl unsubstituted or substituted with 1-3 groups of Ra-

Ra represents C1-6 alkoxy, C1-6 alkyl, CF3, nitro, amino, cyano, C1-6 alkylamino, or halogen;

Rb independently represents H, halogen, C1-6 alkyl, C3-6 cylcoalkyl or

--- represents a double or single bond;

n represents 0-4; and

m represents 0-8.

18(Original). The compound according to claim 17 wherein X and Y are  $(CH_2)_n$ , --- represents a double bond; and Ar<sub>2</sub> is phenyl.

19(Original). The compound according to claim 18 wherein X is (CH<sub>2</sub>)<sub>n</sub> and n is 1 and Y is  $(CH_2)_n$  and n is 3.

#### 20 Cancel.

21(Currently Amended) A pharmaceutical composition for treating ocular hypertension or glaucoma comprising a therapeutically effective amount of a compound of formula I of claim 17 as defined in any one of claims 1 to 10, or a pharmaceutically acceptable salt, enantiomer, diasteromer, prodrug, or mixture thereof, in association with a pharmaceutically acceptable carrier.

22 (Currently Amended) A composition according to claim 21 in a form for topical administration as a solution or suspension and further comprising a second active ingredient as defined in claim 12 or 13.

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- 23 Cancel.
- 24 Cancel.
- 25 Cancel.
- 26 Cancel.
- 27 (New) A composition according to claim 22 wherein a second active ingredient belonging to the group consisting of: b-adrenergic blocking agent, parasympatho-memetic agent, sympathomimetic agent, carbonic anhydrase inhibitor, Maxi-K channel blocker, a prostaglandin, hypotensive lipid, neuroprotectant, and 5-HT2 receptor agonist is added.
- 28 (New) A composition according to claim 27 wherein the βadrenergic blocking agent is timolol, betaxolol, levobetaxolol, carteolol, or levobunolol; the parasympathomimetic agent is pilocarpine; the sympathomimetic agent is epinephrine, brimonidine, iopidine, clonidine, or para-aminoclonidine, the carbonic anhydrase inhibitor is dorzolamide, acetazolamide, metazolamide or brinzolamide; the prostaglandin is latanoprost, travaprost, unoprostone, rescula, or \$1033, the hypotensive lipid is lumigan, the neuroprotectant is eliprodil, R-eliprodil or memantine; and the 5-HT2 receptor agonist is 1-(2-aminopropyl)-3methyl-1H-imdazol-6-ol fumarate or 2-(3-chloro-6-methoxy-indazol-1yl)-1-methyl-ethylamine.